=> d bib abs hitstr ll4 tot L14 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2007 ACS on STN 2005:259680 HCAPLUS AN DN 142:336356 ΤI Preparation of benzimidazoles and imidazopyridines having affinity for melanocortin (MC), in particular MC4, receptors Poitout, Lydie; Brault, Valerie; Sackur, Carole; Roubert, Pierre; Plas, TN Pascale PA U.S. Pat. Appl. Publ., 213 pp., Cont.-in-part of U.S. Ser. No. 504,033. SO CODEN: USXXCO DT Patent English LA FAN.CNT 2 DATE KIND DATE APPLICATION NO. PATENT NO. ---------2004US-0915920 20040811 рΤ US2005065179 A1 20050324 2003FR-0002320 20030226 20040827 FR---2851563 A1 FR---2851563 В1 20050422 2004WO-FR00418 20040225 WO2004075823 A2 20040910 WO2004075823 A3 20041007 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG PRAI 2003FR-0002320 2003US-0504033 20030226 Α A2 20030920 20040225 2004WO-FR00418 W MARPAT 142:336356 os

THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Title compds. I [wherein A = CH2, CO, (un) substituted COCH2; X = CH, N; R1, R2 = independently H, alkyl optionally substituted by OH, alkenyl, etc.; or R1NR2 = (un) substituted hetero(bi) cycloalkyl; R3 = alkyl, alkoxy, alkylthio, heteroaryl, (un) substituted hetero/cycloalkyl, aryl, etc.; R4 = (CH2)sR5; R5 = heterocycloalkyl, heteroaryl, etc.; s = 0-6] were prepared as melanocortin (MC), in particular MC4, receptor modulators (no data given). For example, II was prepared, in 2 steps, by amination of 3-Fluoro-N,N-bis(3-methylbutyl)-4-nitrobenzamide (preparation given) with 3-(piperidino)propylamine in CH3CN at reflux, followed by one-step hydrogenation/coupling with 4-acetylphenyl isothiocyanate. I are useful in the treatment of pathol. states and the diseases in which one or more melanocortin receptors are included such as pain, inflammatory conditions, etc.

IT 848577-67-3P

GT

RE.CNT 10

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of benzimidazoles and imidazopyridines having affinity for melanocortin (MC), in particular MC4, receptors)

RN 848577-67-3 HCAPLUS

CN 1H-Benzimidazole-6-carboxamide, N,N-bis(3-methylbutyl)-1-[3-(1piperidinyl)propyl]-2-[[4-(1-piperidinylsulfonyl)phenyl]amino]- (9CI) (CA
INDEX NAME)

$$Me_2CH - CH_2 - CH_2$$
 $Me_2CH - CH_2 - CH_$

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L14 ANSWER 2 OF 4 HCAPLUS COPYRIGHT 2007 ACS on STN
     2004:817883 HCAPLUS
AN
DN
     141:332190
     Preparation of fused azoles such as 2,5-disubstituted benzimidazoles,
TI
     benzoxazoles and benzothiazoles as kinase inhibitors
     Dipietro, Lucian V.; Harmange, Jean-Christophe; Askew, Benny C., Jr.;
IN
     Elbaum, Daniel; Germain, Julie; Habgood, Gregory J.; Kim, Joseph L.; Patel, Vinod F.; Potashman, Michele; Van der Plas, Simon
PA
     Amgen Inc., USA
     PCT Int. Appl., 289 pp.
     CODEN: PIXXD2
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     Patent
     English
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		CN,	CO,	CR,	CU,	CZ,	DΕ,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,	
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		SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	
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